# IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

PATENT

Applicant(s): Yoshitaka N

Yoshitaka Nishio, et al. Docket No.:

49288.3500

Serial No.:

TBA

Group Art Unit

TBA

Filed:

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Title:

SUBSTRATE DIVIDING SYSTEM, SUBSTRATE Confirmation No.: TBA

MANUFACTURING EQUIPMENT, SUBSTRATE SCRIBING METHOD AND

SUBSTRATE DIVIDING

# INFORMATION DISCLOSURE STATEMENT

Commissioner For Patents Mail Stop Amendment PO Box 1450 Alexandria, VA 22313-1450

### Commissioner:

In accordance with the duty of disclosure under 37 C.F.R. §1.56 and pursuant to 37 C.F.R., §\$1.97 and 1.98, Applicants hereby notify the U.S. Patent and Trademark Office of the documents listed on the attached Form PTO/SB/08A. Applicants respectfully submit that all pending claims are patentable over the foregoing references, alone or in combination. The Examiner is requested to initial the enclosed Form PTO/SB/08A and return a copy thereof to the undersigned.

The submission of the listed documents is not intended as an admission that any such document constitutes prior art against the claims of the present application. Applicants reserve the right to dispute any of the listed documents as prior art during examination. Further, Applicants do not waive any right to take any action that would be appropriate to antedate or otherwise remove any listed document as a competent reference against the claims of the present application. Furthermore, the submission of

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this Information Disclosure Statement is not to be construed as a representation that a search has been made or that no other material information may exist.

- [X] For each of the following items listed on the enclosed copy of Form PTO/SB/08A that is not in the English language, an English language translation of that item or a portion thereof or a concise explanation of the relevance of that item is enclosed.
- 3. [ X ] No fee is due under 37 C.F.R. §1.17(p) for this Information Disclosure Statement since it is being filed in compliance with:
  - [X] 37 C.F.R. §1.97(b)(1), within three months of the filing date of the aboveidentified application.
  - 37 C.F.R. §1.97(b)(2), within three months of the date of entry into the national stage as set forth in §1.491 in an international application.
  - 37 C.F.R. §1.97(b)(3), before the mailing date of a first Office action on the merits.
- 4. [] No fee is due under 37 C.F.R. §1.17(p) for this Information Disclosure Statement since it is being filled in compliance with 37 C.F.R. §1.97(c), after the period specified in paragraph 3 above but before the mailing date of a final action or a Notice of Allowance (where there has been no prior final action), and is accompanied by one of the certifications pursuant to 37 C.F.R. §1.97(e) set forth in paragraph 8 below.
- 5. [] A fee is due under 37 C.F.R. §1.17(p) for this Information Disclosure Statement since it is being fled in compliance with 37 C.F.R. §1.97(c), after the period specified in paragraph 3 above but before the mailing date of a final action or a notice of allowance (where there has been no prior final action):
  - [ ] A check in the amount of \$180.00 is enclosed in payment of the fee.
  - Charge the fee to Deposit Account No. 19-2814.
- 6. [] A fee is due under 37 C.F.R. §1.17(i)(1) for this Information Disclosure Statement since it is being filed in compliance with 37 C.F.R. §1.97(d), after the mailing date of a final action or a notice of allowance, whichever comes first, but before payment of the issue fee, and is accompanied by:

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- one of the certifications pursuant to 37 C.F.R. §1.97(e) set forth in paragraph 8 below; and
- the attached petition requesting consideration of this Information Disclosure Statement; and
- the fee due under 37 C.F.R. §1.17(i)(1) which is paid as set forth in paragraph 9 below.
- A fee is due under 37 C.F.R. §1.17(i)(1) for this Information Disclosure Statement since it is being filed in compliance with:
  - a. [] 37 C.F.R. §1.313(b)(3), after the issue fee has been paid and information cited in this Information Disclosure Statement may render at least one claim unpatentable and is accompanied by the attached Petition To Withdraw Application From Issue:
  - b. [] 37 C.F.R. §1.313(b)(5), after the issue fee has been paid and information cited in this Information Disclosure Statement is to be considered in a Continuation application upon abandomment of the instant application and is accompanied by the attached Petition To Withdraw Application From Issue.
  - The fee due under 37 C.F.R. §1.17(i)(1) is paid as set forth in paragraph 9 below.
- 8. [] I hereby certify that each item of information contained in this Information Disclosure Statement was cited in a communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of this Information Disclosure Statement.
  - [] I hereby certify that no item of information in the Information Disclosure Statement filed herewith was clated in a communication from a foreign patent office in a counterpart foreign application or, to my knowledge after making reasonable incuiry, was known to any individual designated in §1.56(c) more than three months prior to the filing of this Information Disclosure Statement.
- A check in the amount of \$180.00 is enclosed in payment of the fee due under 37 C.F.R. §1.17(p).
  - [ ] Charge the fee due under 37 C.F.R. §1.17(i)(1) to Deposit Account No. 19-2814.

[ X ] The Commissioner is hereby authorized to charge any additional fees which may be required for this Information Disclosure Statement, or credit any overpayment to Deposit Account No. 19-2814.

Respectfully submitted,

Registration No. 39.038

Snell & XV

d: 9/13/06

Snell & Wilmer L.L.P. One Arizona Center 400 E. Van Buren

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| Application Number Filing Date          |                                 |                      |                                       |
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|   | cation Number                   | Application Number   |                                       |
|   | Date                            | Filing Date          | INFORMATION DIGGS COURT               |
| STATEMENT BY APPLICANT                  | Named Inventor Yoshitaka Nishio | First Named Inventor | INFORMATION DISCLOSURE                |
| ( Not for submission under 37 CFR 1.99) | nit TBA                         | Art Unit             |                                       |
| Examiner Name TBA                       | niner Name TBA                  | Examiner Name        | (Notice submission under or or it may |
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|                      | 1           | 10-209086                               | JP                           |                 |          | 1998-08-07                                      | Matsushita Electric<br>Co Ltd.                        | Ind   |          |             |    |
|                      | 2           | 2000-264657                             | JP                           |                 |          | 2000-09-26                                      | Mitsuboshi Diamor<br>Kogyo KK                         | nd  |          |             |    |
|                      | 3           | 2001-347497                             | JP                           |                 |          | 2001-12-18                                      | Hitachi Ltd   |   |          |             |    |

|   | Application Number        |  |              |
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| NFORMATION DISCLOSURE<br>STATEMENT BY APPLICANT | First Named Inventor Yosh |  | itaka Nishio |
| Not for submission under 37 CFR 1.99)           | Art Unit                  |  | TBA          |
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### CERTIFICATION STATEMENT

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 37 CFR 1.97(e)(1).

## OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquiry, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 1.59(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 1.39(file).

- 7 See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.

Please see 37 CFR 1 97 and 1 98 to make the appropriate selection(s):

☐ None

#### SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

| Torn or the agricult. |            |                     |                     |            |  |  |  |
|-----------------------|------------|---------------------|---------------------|------------|--|--|--|
|                       | Signature  | /Howard I Sobelman/ | Date (YYYY-MM-DD)   | 2006-09-13 |  |  |  |
|                       | Name/Print | Howard I Sobelman   | Registration Number | 39038      |  |  |  |

This collection of information is required by 37 CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is to file fand by the USPTO to process) an application. Confidentially is governed by \$5 U.S. C.12 and 37 CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patient and Tradenant's Office, U.S. operationed for Commence, P. 0. Bot 1450, Alexandria, V.S. 2213.1-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.2.2313.1-1450.

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The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kollie is to process and/or examine your submission related to a patient application or patient. If you do not furnish the requested process and/or examine your submission related to a patient application or patient. If you do not furnish the requested related to the patient of the principal principal

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
  - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiation.
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  - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
    may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
    to the Patent Cooperation Treaty.
  - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
- 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designe, during an inspection of records conducted by GSA a part of that agency's responsibility to recommend improvements in records management practices and programs, under suthority of 4d U.S.C. 2004 and 2006. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant for the control individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 12(b) or issuance of a patient pursuant to 35 U.S. C. 15.1 Further, a record may be disclosed, subject to the limitations of 37 CFR.114, as a routine use, to the public if the record via flori of mapplication which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inseptions or an issued patient.
  - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.